Docket No.: B235 1010.1 Application No.: 10/748,094

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

1. (currently amended) A process for manufacture of long circulating non-pegylated liposomes comprising:

dissolving one or more phospholipids and one or more sterols in a solvent or mixture of solvents;

hydrating the phospholipids and sterols with an aqueous hydration media; removing the solvent or mixture of solvents before or after hydrating the lipids <u>and sterols</u>;

sizing the non-pegylated liposomes formed to about 0.06 μm to form a liposomal composition;

removing extraliposomal hydration salt from the liposomal composition using a sucrosehistidine buffer solution;

wherein the amount of aqueous hydration media used is in the range of 10 to 35 ml for each mmole of phospholipid present to form non-pegylated liposomes; and

wherein the aqueous hydration media comprises ammonium sulfate and sucrose; and wherein the one or more phospholipids is a saturated phosphatidylcholine selected from the group consisting of distearoyl phosphatidylcholine (DSPC), dipalmitoyl phosphatidylcholine (DPPC), hydrogenated soya phosphatidylcholine (HSPC) and derivatives mixtures thereof; and wherein the one or more phospholipids exhibits a phase transition temperature of between 50 and 65°C; and

wherein the <u>resulting</u> non-pegylated liposomes have a blood circulation half life of at least 25 times longer than conventional non-liposomal formulations when tested in Swiss albino mice at equivalent doses.

2. (original) The process of claim 1 wherein the amount of aqueous hydration media used is 30 ml for each mmole of phospholipid in the lipid solution.

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3. (original) The process of manufacture of non-pegylated liposomes of claim 1 further

comprising loading the liposomes with a therapeutic or diagnostic agent.

4. (original) The process of claim 3, wherein the therapeutic agent is an antineoplastic

agent.

5. (original) The process of claim 4, wherein the antineoplastic agent is selected from the

group consisting of Doxorubicin hydrochloride, Daunorubicin hydrochloride, and Epirubicin

hydrochloride.

6. (original) The process of claim 5, wherein the antineoplastic agent is Doxorubicin

hydrochloride.

7. (original) The process of claim 1, wherein the molar ratio of phospholipid to sterol is

from about 1:0.1-1:2.

8. (currently amended) The process of claim 7, wherein the wherein the molar ratio of

phospholipid to sterol is from about 1:0.7.

9. (previously canceled).

10. (previously presented) The process of claim 1, wherein the concentration of ammonium

sulfate in aqueous hydration media is not less than 125 mmoles/liter.

11. (previously canceled).

12. (previously presented) The process of claim 1, wherein the phospholipid has a minimum

of sixteen carbons fatty acid chain.

13. (previously canceled).

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14. (previously presented) The process of claim 1, wherein the phospholipid is distearoyl

phosphatidylcholine (DSPC) and wherein the sterol is cholesterol.

15. (original) The process of claim 1, wherein the non-pegylated liposomes are successively

extruded through series of filters having pore sizes from 0.4 µm to 0.05 µm for sizing.

16. (original) A liposome manufactured by the process of claim 1.

17. (original) The liposome of claim 16, wherein the phospholipid comprises distearoyl

phosphatidylcholine (DSPC) and the sterol comprises cholesterol.

18. (Original) The liposome of claim 16, wherein the non-pegylated liposome further

comprises a therapeutic or diagnostic agent.

19. (Original) The liposome of claim 18, wherein said therapeutic agent comprises an

antineoplastic agent.

20. (original) The liposome of claim 19, wherein the antineoplastic agent is selected from the

group consisting of Doxorubicin hydrochloride, Daunorubicin hydrochloride, and Epirubicin

hydrochloride.

21. (original) The liposome of claim 20, wherein the antineoplastic agent is Doxorubicin

hydrochloride.

22. (original) The liposome of claim 16, wherein the average size of liposome is 0.06 µm to

0.16 µm in diameter.

23-61. (previously canceled).

62. (Currently canceled).

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